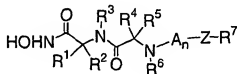


1. A compound selected from the group of compounds represented by Formula (I):



5 wherein:

R² is: (i) cycloalkyl, cycloalkylalkyl, aryl, aralkyl, aralkenyl, heteroaryl, heteroaralkyl, aralkenyl, heterocyclo or heterocycloalkyl; or

(iii) -(alkylene)-B-X where B is -NR⁸CO- (where R⁸ is H or alkyl), and X is cycloalkyl, cycloalkylalkyl, aryl, aralkyl heteroaryl or heteroaralkyl; or

15 R^3 is hydrogen or alkyl;

 R^5 is:

(ii) R^5 and R^4 form an alkylene chain; or

n is 0 or 1;

A is $-C(=O)-CH(R^9)-(CH_2)_m-N(R^{10})-$ wherein:

m is an integer from 0-5 inclusive;

R⁹ is hydrogen, alkyl, cycloalkyl, cycloalkylalkyl, aryl, aralkyl, heteroaryl, heteroaralkyl, heterocycloalkyl, heteroalkyl, or -(alkylene)-C(O)-X¹ where X¹ is alkyl, hydroxy, alkoxy, aryl, aralkyl, aryloxy, aralkyloxy, heteroaryl, heteroaryloxy, heteroaralkyloxy or NR'R''

5 (where R' and R'' are independently H or alkyl, or R' and R'' form an alkylene chain); and

R¹⁰ is hydrogen, alkyl, aralkyl or heteroaralkyl;

Z is Y-B wherein:

Y is alkylene or a bond; and

10 B is -CO-, -C(O)O-, -CONR⁸-, -SO₂-, or -SO₂NR⁸- (where R⁸ is hydrogen or alkyl), alkylene (optionally substituted by hydroxy, alkoxy, amino, monoalkylamino or dialkylamino) or a bond;

R⁷ is cycloalkyl, cycloalkylalkyl, aryl, aralkyl, heteroaryl or heteroaralkyl;

provided that when n = 0 and Z is SO₂, then R² does not contain an imidazole group; and their pharmaceutically acceptable salts, prodrugs, individual isomers, and mixtures of isomers.

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2. The compound of Claim 1 wherein:
n is 0.

3. The compound of Claim 2 wherein R³ and R⁶ are hydrogen.

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4. The compound of Claim 3, wherein:
R² is aralkyl or heteroaralkyl.

5. The compound of Claim 4 wherein:
Z is -C(O)O- or -S(O)₂-.

6. The compound of Claim 5 wherein:
R² is optionally substituted benzyl or heteroaralkylmethyl.

7. The compound of Claim 6 wherein, R^2 is 4-t-butoxybenzyl, 3-chlorobenzyl, 3-indolyl methyl, 2-thienylmethyl, 4-imidazolymethyl or 4-thiazolymethyl.

8. The compound of Claim 7 wherein R^2 is 4-thiazolymethyl.

9. The compound of Claim 8 wherein:

R^7 is aryl, aralkyl, heteroaryl or heteroaralkyl.

10. The compound of Claim 8 wherein:

Z is $-C(O)O-$ and R^1 is optionally substituted benzyl.

11. The compound of Claim 9 wherein:

Z is $-SO_2-$ and R^7 is aryl or heteroaryl.

12. The compound of Claim 10, wherein:

R^1 and R^4 are hydrogen and R^5 is alkyl.

13. The compound of Claim 12 wherein R^5 is (S,S)-1-methylpropyl.

14. The compound of Claim 11, wherein:

R^1 and R^4 are hydrogen and R^5 is alkyl.

15. The compound of Claim 14 wherein R^5 is (S,S)-1-methylpropyl.

16. The compound of Claim 3 wherein:

R^2 is (alkylene)-B-X where B is $-O-$, $-NR^8$, $-S(O)_n-$ (where n is 0, 1 or 2), $-C=O$, $-CONR^8$, $-NR^8CO_2-$, $-NR^8SO_2-$ or $-C(=NR^8)NSO_2-$ (where R^8 is H or alkyl), and X is cycloalkyl, cycloalkylalkyl, aryl, aralkyl heteroaryl or heteroaralkyl.

17. The compound of Claim 16, wherein:

Z is -C(O)O- or -S(O)₂-.

18. The compound of Claim 17, wherein R² is CH₂-B-X and

B is -NHCO₂- and X is benzyl.

19. The compound of Claim 18 wherein:

R⁷ is aryl or aralkyl.

20. The compound of Claim 19, wherein:

R¹ and R⁴ are hydrogen and R⁵ is alkyl.

21. The compound of Claim 20 wherein R² is (S,S)-1-methylpropyl.

22. The compound of Claim 1 wherein:

n is 1.

23. The compound of Claim 22 wherein m is 0 and R³ and R⁶ are hydrogen.

24. The compound of Claim 23, wherein:

R² is aralkyl or heteroaralkyl.

25. The compound of Claim 24, wherein:

Z is -C(O)O- or -S(O)₂-.

26. The compound of Claim 25, wherein:

R² is optionally substituted benzyl or heteroaralkylmethyl.

27. The compound of Claim 26 wherein R² is 4-t-butoxybenzyl, 3-chlorobenzyl, 3-indolyl methyl, 2-thienylmethyl, 4-imidazolylmethyl or 4-thiazolylmethyl.

28. The compound of Claim 27 wherein R^2 is 4-thiazolylmethyl.

29. The compound of Claim 28 wherein:
 R^7 is aryl, aralkyl, heteroaryl or heteroaralkyl.

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30. The compound of Claim 29 wherein:
 Z is $-C(O)O-$ and R^7 is benzyl.

31. The compound of Claim 29 wherein:
 Z is $-SO_2-$ and R^7 is aryl.

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32. The compound of Claim 30, wherein:
 R^1 and R^4 are hydrogen and R^5 is alkyl.

15 33. The compound of Claim 32 wherein R^5 is (S,S)-1-methylpropyl.

34. The compound of Claim 31, wherein:
 R^1 and R^4 are hydrogen and R^5 is alkyl.

20 35. The compound of Claim 34 wherein R^5 is (S,S)-1-methylpropyl.

36. The compound of Claim 23, wherein:
 R^2 is (alkylene)-B-X where B is $-O-$, $-NR^8$, $-S-$, $-C=O$, $-CONR^8$, $-NR^8CO_2$, $-NSO_2$ or $-C(=NR^8)NSO_2$ (where R^8 is H or alkyl), and X is cycloalkyl, cycloalkylalkyl, aryl, aralkyl, heteroaryl or heteroaralkyl.

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37. The compound of Claim 36, wherein:
 Z is $-C(O)O-$ or $-S(O)_2-$.

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38. The compound of Claim 37, wherein R^2 is CH_2-B-X and B is $-NHCO_2-$ and X is benzyl.

39. The compound of Claim 38 wherein:

5 R^7 is aryl or aralkyl.

40. The compound of Claim 39, wherein:

R^1 and R^4 are hydrogen and R^5 is alkyl.

10 41. The compound of Claim 40 wherein R^5 is (S,S)-1-methylpropyl.

42. A pharmaceutical composition comprising the compound of Claim 1 and a pharmaceutically acceptable excipient.

15 43. A method of treating disease comprising administering to a patient in need thereof a compound of Claim 1.

44. The method of Claim 43, wherein the disease is a fibrotic disease.

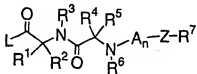
20 45. The method of Claim 44 wherein the disease is acute respiratory distress syndrome.

46. A method of treating fibrosis comprising administering to a patient in need thereof an inhibitor of procollagen C-proteinase that is at least ten-fold more selective for procollagen C-proteinase over both collagenase-1, collagenase-2 and collagenase-3.

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47. A method for preparing the compounds of Claim 1 by:
- (i) treating a compound of Formula II wherein L is a leaving group and $R^1 - R^7$, A, n and Z are as defined in Claim 1 with hydroxylamine or a protected derivative thereof, and
 - (ii) deprotecting as necessary and isolating the compound of Claim 1.

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(II)

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